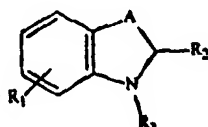
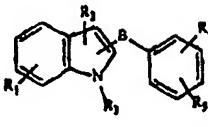
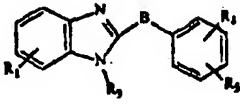




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : C07D 209/12, A61K 31/40, C07D 209/42, 401/12, 209/10, 405/12		A1	(11) International Publication Number: WO 98/08818
			(43) International Publication Date: 5 March 1998 (05.03.98)
(21) International Application Number: PCT/US97/14943 (22) International Filing Date: 26 August 1997 (26.08.97) (30) Priority Data: 08/703,115 26 August 1996 (26.08.96) US (71) Applicant: GENETICS INSTITUTE, INC. [US/US]; 87 CambridgePark Drive, Cambridge, MA 02140 (US). (72) Inventors: XIANG, YiBin; 821 Main Street, Acton, MA 01720 (US). BEMIS, Jean; 256 Appleton Street, Arlington, MA 02174 (US). MCKEW, John; 58 Varnum Street, Arlington, MA 02174 (US). KAILA, Neelu; 2 Course Brook Lane, Natick, MA 01760 (US). (74) Agent: BROWN, Scott, A.; Genetics Institute, Inc., 87 CambridgePark Drive, Cambridge, MA 02140 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>	
(54) Title: INHIBITORS OF PHOSPHOLIPASE ENZYMES			
(57) Abstract			
<p>Compounds having a chemical formula selected from the group consisting of formulae (I), (II) and (III) or a pharmaceutically acceptable salt thereof, wherein: A is independent of any other group and is selected from the group consisting of -CH₂- and -CH₂-CH₂-; B is independent of any other group and is selected from the group consisting of -(CH₂)_n-, -(CH₂O)_n-, -(CH₂S)_n-, -(OCH₂)_n-, -(SCH₂)_n-, -(CH=CH)_n-, -(C≡C)_n-, -CON(R₆)-, -N(R₆)CO-, -O-, -S- and -N(R₆)-; R₂ is independent of any other R group and is selected from the group consisting of -H, -COOH, -COR₅, -CONR₅R₆, -(CH₂)_n-W-(CH₂)_m-Z-R₅, -(CH₂)_n-W-R₅, -Z-R₅, C₁-C₁₀ alkyl, alkenyl and substituted aryl; R₃ is independent of any other R group and is selected from the group consisting of -H, -COOH, -COR₅, -CONR₅R₆, -(CH₂)_n-W-(CH₂)_m-Z-R₅, -(CH₂)_n-W-R₅, -Z-R₅, C₁-C₁₀ alkyl, alkenyl and substituted aryl; which inhibit the activity of phospholipase enzymes, particularly cytosolic phospholipase A₂. Pharmaceutical compositions comprising such compounds and methods of treatment using such compositions are also disclosed.</p>			
<div style="display: flex; justify-content: space-around; align-items: center;"> <div style="text-align: center;">  <p>(I)</p> </div> <div style="text-align: center;">  <p>(II)</p> </div> </div> <div style="text-align: center; margin-top: 20px;">  <p>(III)</p> </div>			